CVS LUBRICANT EYE DROPS- carboxymethylcellulose sodium liquid
CVS Pharmacy, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Lubricant

Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eyes or exposure to wind or sun
- may be used as a protectant against further irritation

Warnings

For external use only

Do not use this product if

- solution changes color or becomes cloudy

When using this product

- do not reuse
- once opened, discard
- to avoid contamination, do not touch tip of container to any surface
- do not touch unit-dose tip to eye

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye worsens or persists for more than 72 hours

Keep out of the reach of children.

If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- to open, twist and pull tab to remove
- instill 1 or 2 drops in the affected eye(s) as needed and discard container
- if used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions

Other information

- store at 15°-25°C (59°-77°F)
- use only if single-use container is intact
- use before expiration date marked on container
- RETAIN THIS CARTON FOR FUTURE REFERENCE

Inactive ingredients

calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid to adjust pH.

CVS Lubricant Eye Drops Preservative Free 70 ct

CVS LUBRICANT EYE DROPS

carboxymethylcellulose sodium liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-804
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-804-01	70 in 1 BOX	07/30/2019	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	07/30/2019	

Labeler - CVS Pharmacy, Inc. (062312574)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(69842-804) , label(69842-804)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(69842-804)

Revised: 7/2019

CVS Pharmacy, Inc.